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PTO/SB/05 (12/97) (modified)
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06/12/98

UTILITY PATENT APPLICATION TRANSMITTAL <small>(Only for new nonprovisional applications under 37 CFR 1.53(b))</small>	Attorney Docket No	290252021800	Total Pages	3
	First Named Inventor or Application Identifier			
	Jill McFadden et al.			
	Express Mail Label No	EL053907928US		

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<p>1. <input checked="" type="checkbox"/> Fee Transmittal Form <small>(Submit an original, and a duplicate for fee processing)</small></p> <p>2. <input checked="" type="checkbox"/> Specification <small>[Total Pages 18]</small> <small>(preferred arrangement set forth below)</small></p> <ul style="list-style-type: none">- Descriptive title of the Invention- Cross References to Related Applications- Statement Regarding Fed sponsored R & D- Reference to Microfiche Appendix- Background of the Invention- Brief Summary of the Invention- Brief Description of the Drawings <small>(if filed)</small>- Detailed Description- Claim(s)- Abstract of the Disclosure <p>3. <input checked="" type="checkbox"/> Drawing(s) <small>(35 USC 113)</small> <small>[Total Sheets 2]</small></p> <p>4. Oath or Declaration <small>[Total Pages 2]</small></p> <p>a. <input checked="" type="checkbox"/> Newly unexecuted (original or copy)</p> <p>b. <input type="checkbox"/> Copy from a prior application (37 CFR 1.63(d)) <small>(for continuation/divisional with Box 17 completed)</small> [Note Box 5 below]</p> <p>c. <input type="checkbox"/> DELETION OF INVENTOR(S) <small>Signed statement attached deleting inventor(s) named in the prior application, see 37 CFR 1.63(d)(2) and 1.33(b)</small></p> <p>5. <input type="checkbox"/> Incorporation By Reference <small>(useable if Box 4b is checked)</small> <small>The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 4b, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein</small></p>	<p>6. <input type="checkbox"/> Microfiche Computer Program <small>(Appendix)</small></p> <p>7. Nucleotide and/or Amino Acid Sequence Submission <small>(if applicable, all necessary)</small></p> <p>a. <input type="checkbox"/> Computer Readable Copy</p> <p>b. <input type="checkbox"/> Paper Copy (identical to computer copy)</p> <p>c. <input type="checkbox"/> Statement verifying identity of above copies</p>
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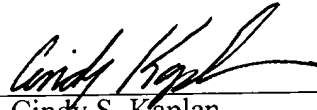
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Dated: June 12, 1998

Respectfully submitted,

By: _____



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5 *June 12, 1998* *Krista K. Merrimac*
Date Krista K. Merrimac

CATHETER WITH KNIT SECTIONFIELD OF THE INVENTION

10 This invention relates generally to catheters, and more particularly, to a catheter having a knit section.

BACKGROUND OF THE INVENTION

15 Catheters are increasingly used to access remote regions of the human body and, in doing so, delivering diagnostic or therapeutic agents to those sites. In particular, catheters which use the circulatory system as the pathway to these treatment sites are especially practical. Catheters are also used to access other regions of the body, e.g., genito-urinary regions, for a variety of therapeutic and diagnostic reasons. One such treatment of diseases of the circulatory system is
20 via angioplasty (PCA). Such a procedure uses catheters having balloons on their distal tips. It is similarly common that those catheters are used to deliver a radiopaque agent to the site in question prior to the PCA procedure to view the problem prior to treatment.

25 Often the target which one desires to access by catheter is within a soft tissue such as the liver or the brain. These are difficult sites to reach. The catheter must be introduced through a large artery such as those found in the groin or in the neck and then be passed through ever-narrower regions of the arterial system until the catheter reaches the selected site. Often such pathways will wind back upon themselves in a multi-looped path. These catheters are difficult to
30 design and to utilize in that they must be fairly stiff at their proximal end so to allow the pushing and manipulation of the catheter as it progresses through the body, and yet must be sufficiently flexible at the distal end to allow passage of the catheter tip through the loops and increasingly smaller blood vessels mentioned

above and yet at the same time not cause significant trauma to the blood vessel or to the surrounding tissue. Further details on the problems and an early, but yet effective, way of designing a catheter for such a traversal may be found in U.S. Patent No. 4,739,768, to Engelson. U.S. Patent No. 5,702,373 discloses another catheter design for use in accessing a tissue target within the body. The catheter includes a braided reinforcing member for high kink resistance.

These catheters are designed to be used with a guidewire. A guidewire is simply a wire, typically of very sophisticated design, which is the "scout" for the catheter. The catheter fits over and slides along the guidewire as it passes through the vasculature. Said another way, the guidewire is used to select the proper path through the vasculature with the urging of the attending physician and the catheter slides along behind once the proper path is established.

There are other ways of causing a catheter to proceed through the human vasculature to a selected site, but a guidewire-aided catheter is considered to be both quite quick and somewhat more accurate than the other procedures. One such alternative procedure is the use of a flow-directed catheter. These devices often have a small balloon situated on the distal end of the catheter which may be alternately deflated and inflated as the need to select a route for the catheter is encountered.

This invention is an adaptable one and may be used in a variety of catheter formats. The invention utilizes the concept of combining one or polymeric tubes with a knit tube. The construction technique has the benefit of producing catheter sections having small overall diameters but with exceptional strength, resistance to ovalization and kinking, and recovery from kinking (even in vivo) should such kinking occur. The catheter may be used in conjunction with a guidewire, but the catheter may also be used as a flow-directed catheter with the attachment of a balloon or in combination with a specifically flexible tip.

Knit tubular members have been used in surgical devices such as stents. For example, U.S. Patent No. 5,662,713 discloses a self-expanding stent formed by knitting a wire into a pattern of overlapping loops. The stent is placed within the body to perform a function such as maintaining open a body lumen. U.S.

Patent No. 5,480,423 discloses a self-expanding knit stent and a catheter for delivering the stent.

None of the documents cited above provides a catheter structure described in the specification and set forth in the claims recited below, particularly when the flexibility and ability to resist kinks is factored into the physical description of the devices.

SUMMARY OF THE INVENTION

This invention is a catheter section having a knit tubular member. The catheter section may include an inner tubular liner in coaxial relationship with the knit tubular member and an outer tubular cover surrounding the knit tubular member. The knit tubular member may comprise a metal alloy, polymeric material or other suitable material.

The catheter section may be used as a distal segment of a catheter with a relatively more stiff proximal segment. The proximal segment may also include an inner liner and outer cover. A stiffener formed from a knit member, braid, or coil may also be interposed between the inner liner and outer cover.

The kink resistance of the catheter is due to the presence of the knit tubular member. In addition to exceptional kink resistance, the catheter section may be made in such a way that the wall is extraordinarily thin, particularly when compared to walls of catheters having equal strength but made solely of polymeric materials. The catheter section additionally is very resilient in that, should the catheter section be kinked, the kink is self-healing. This resiliency means that the catheter need not be withdrawn from a patient's vasculature simply because the catheter has inadvertently kinked. Simple movement of the catheter will cure the kink. Kinking minimization is a matter of concern with many catheters in the marketplace today. The knit tubular member also minimizes circumferential collapse (ovaling of the catheter section) and increases the burst strength of the catheter section.

The catheter assembly may also have ancillary components such as a luer lock and some manner of providing radiopacity.

The above is a brief description of some deficiencies in the prior art and advantages of the present invention. Other features, advantages, and embodiments of the invention will be apparent to those skilled in the art from the following description, drawings, and claims.

5

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a side view of a typical three section catheter made using the concepts of this invention;

10

Fig. 2 is an enlarged view of a portion of the catheter of Fig. 1;

Fig. 3 is a side view of a distal section of the catheter of Fig. 1; and

Fig. 4 is an enlarged partial view of a knit tubular member of the catheter of Fig. 1.

15

Corresponding reference characters indicate corresponding parts throughout the several views of the drawings.

DESCRIPTION OF THE INVENTION

20

This invention is a catheter assembly having a more distal segment preferably made up of an inner liner, an outer covering, and a knitted layer located between the liner and the covering, and a more proximal segment comprising, for example, a stiff polymeric or metallic tubing member. Other sections of these or other designs may be placed variously between the noted sections or distal of the distal knitted section noted above.

25

A typical multi-section catheter 100 which may incorporate the concepts of this invention is shown in Figure 1. Such a catheter is described in more detail in U.S. Patent No. 4,739,768, to Engelson, the entirety of which is incorporated by reference and is particularly suitable for neurological and peripheral vascular applications. Clearly, then, it is also suitable for less demanding service such as might be encountered in access and treatment of the heart. One difficulty which has arisen as higher demands for length have been placed on these catheters is that the diameter of the distal section necessarily becomes smaller and smaller. This is

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so since the longer catheters must reach ever smaller vascular areas. This smaller diameter requires a concomitant thinning of the wall section. The thinner section walls may kink or ripple when actively pushed along the guidewire or when vaso-occlusive devices are pushed through the catheter's lumen. The typical configuration shown in Figure 1 has a distal section 102 having significant flexibility, an intermediate section 104 which is typically less flexible, and a long proximal section 106 which in turn is least flexible. The distal section 102 is flexible and soft to allow deep penetration of the extraordinary convolutions of the neurological vasculature without trauma. Various known and often necessary accessories to the catheter assembly, e.g., one or more radiopaque bands 108 at the distal region to allow viewing of the position of the distal region under fluoroscopy and a luer assembly 110 for guidewire 112 and fluids access, are also shown in Figure 1. The typical dimensions of this catheter are:

Overall length:	60-200 cm
Proximal Section (106):	30-150 cm
Intermediate Section (104):	20-50 cm
Distal Section (102):	2.5-30 cm

Obviously, these dimensions are not particularly critical to this invention and are selected as a function of the malady treated and its site within the body. Typical of the catheters made using this invention are those in the 2 French to 8 French range (1 French = 0.013 inch). The inner diameter of such catheters is then 10 mils. to 91 mils. (1 mil = .001 inch).

Furthermore, a catheter made using this inventive concept need not be of three sections with increasing stiffness as is shown in Figure 1. The catheter may be of two discrete sections or may be of four or more discrete sections of differing flexibility. Through judicious choice of physical parameters for the catheter sections, the components may also have varying physical parameters (e.g., lubricity, flexibility, wall thickness, inner or outer layer member composition, etc.) within the sections. Typically, although not necessarily, when a three section

catheter is desired, the most proximal section 106 is the "more proximal" or "stiff" section described herein. Again, although not necessarily, when a three section catheter is desired, the most distal section 102 is the "more distal" or "flexible" (soft) section.

5 Fig. 2 shows one variation of a distal segment 122 having a knit tubular member 128, and a more proximal segment 124 of a catheter. As previously discussed, the more proximal segment 124 is stiffer than the distal segment 122. The knit tubular member 128 may also extend into the more proximal segment 124.

10 Fig. 3 shows an enlarged view of the more distal segment 122 with an outer tubular cover 130 extending over the knit tubular member 128. The knit member 128 is interposed between the outer cover 130 and an inner tubular liner 126. Both the inner liner 126 and the outer cover 130 are preferably polymeric and may be melt-miscible. The materials of the inner and outer layers 126, 130
15 may also contain adhesives. The outer cover 130 is preferably heat-shrunk, extruded, or bonded onto the inner liner 126 and knit member 128 as further described below.

 Preferred polymeric materials for the inner liner 126 include fluoropolymers including PTFE, FEP, vinylidene fluoride, polyethylene,
20 polyvinyl chloride (PVC), ethyl vinyl acetate (EVA), polyethylene terephthalate (PET), Pebax (polyether block amide), Nylon, and their mixtures, alloys, copolymers and block copolymers. Another useful class of polymers are thermoplastic elastomers, including those containing polyesters as components. Typical of this class is HYTREL. The inner liner 126 may be coated onto the
25 inner surface of the knit member 128. Polyesters and polyimides, in particular, are useful as adhesives in this service. The inner liner 126 may also be formed by dipping a mandrel into a molten polymer, for example. The wall thickness of the inner liner 126 may be between 0.3 mils. and 3.0 mils., for example.

 The outer cover 130 may be made of polyethylene or of EVA or their
30 mixtures, for example. Preferred polymeric materials for the outer cover 130 include polyimide, polyamide, polypropylene, fluoropolymers including PTFE,

FEP, Pebax, Nylon, vinylidene fluoride, and their mixtures, alloys, copolymers, and block copolymers. The polymer is typically extruded onto a tubing of appropriate size and thickness and then crosslinked to raise the melt temperature of the resulting tubing. The tubing is then inflated and perhaps stretched to give the included polymer a specific molecular orientation. The tubing, so treated, may then be slipped over the combination of inner liner 126 and knit member 128 and heat shrunk into place. If a section with even more flexibility is required, the outer cover 130 may also be of a member selected from a more flexible material such as polyurethane, low density polyethylene (LDPE), polyvinylchloride, THV, and other polymers of suitable softness or modulus of elasticity. The wall thickness of the outer cover 130 may be between 0.5 mils. and 10 mils., for example.

It is to be understood that the materials and thickness of the inner liner 126 and outer cover 130 may be different than described herein without departing from the scope of the invention. Furthermore, each of the polymers noted herein may be used in conjunction with radiopaque filler materials such as barium sulfate, bismuth trioxide, bismuth carbonate, powdered tungsten, powdered tantalum, or the like so that the location of various portions of the catheter sections may be radiographically visualized within the human body.

The knit tubular member 128 is preferably formed from a metal alloy, such as stainless steel, tantalum or its alloys, tungsten, platinum, or a superelastic alloy. Preferred superelastic alloys include the class of nickel/titanium materials known as nitinol, disclosed in U.S. Patent Nos. 3,174,851, 3,351,463, and 3,753,700. Commercial alloys containing up to about 5% of one or more other members of the iron group, e.g., Fe, Cr, Co, are considered to be encompassed within the class of superelastic nickel/titanium alloys suitable for use. When using a superelastic alloy, an additional heat treat step may be desirable to preserve the shape of the knit tubular member 128. After heat treatment, the knit member 128 retains its shape and the alloy retains its superelastic properties.

The knit member 128 may also be formed from a non-metal alloy such as carbon/graphite, aramids (e.g., Kevlar, Tehnora, Twaron), liquid crystal polymer

(LCP) (e.g., Vectran, PBO), polyethylene (e.g., Spectra, Dyneema, Certran), polyester (e.g., Dacron, Pentex, Dacronr), or Nylon.

Fig. 4 shows a front half of the knit tubular member 128. The knit member 128 is preferably formed from a wire 134 having a generally circular cross-sectional shape. The diameter of the wire is preferably about 0.5 mil. - 1.5 mil. The loop length L of the knit may be about 3.0 mil. - 15.0 mil., and the peak-to-peak height H of the loops may be about 3.0 mil. - 30.0 mil., for example. The wire may have other cross-sectional shapes such as rectangular (i.e., ribbon), for example. The knit tubular member 128 is preferably knitted from a single strand and configured with the “down loops” and “up loops” having the same size. Multiple strands or other knit configurations may also be used. For example, multiple strands of the same or different material may be knit together (i.e., co-knit), as is well known by those skilled in the art. The knit member 128 may also be formed from a multifilament wire comprising stainless steel, platinum, nitinol, or any combination of these or other materials. For example, the multifilament wire may comprise stainless steel and platinum, or nitinol and platinum.

The knit member 128 is preferably tightly knitted so that it is not significantly radially expandable (e.g., does not increase in diameter more than about 5% when an outwardly directed radial force is applied to an inner surface of the knit member).

The knit member 128 may be knitted on a conventional knitting machine, such as an Emilnestle machine, (product designation number E88-187, or E90-102). The knitting machine produces a long rope of knit loops (Fig. 4). After knitting, the tubular member 128 may be mounted on a mandrel for annealing, to relieve strains induced by the plastic deformation of knitting and to produce greater elasticity in the wire 134, if a superelastic alloy is used, for example. After annealing, the member 128 is cut to its final length. The knit member may also be formed directly on the inner liner 126. To prevent the knit member 128 from unraveling, the last three loops may be coated or the outer cover may be used to prevent unraveling. The sharp ends of the wire 134 are covered by the outer cover 130.

The number and type of sections of the catheter may vary. The catheter sections may have multiple polymeric layers exterior of the knit member 128 as well as multiple polymeric liner members interior to the knit member. The catheter sections may also include multiple layers of knit members 128 between or amongst the various polymer layers. Each individual section of the catheter may also vary in stiffness. Different sections may be connected together by soldering or with an adhesive, for example. An outer cover, as described above, may also be applied to the outer surface of both the more distal segment 122 and the more proximal segment 124 to connect the two sections together. The outer cover of the proximal and distal sections may also be formed as interlocking components. For example, adjacent sides of the outer covers may have opposing interlocking teeth or wedge shaped members which fit together to connect two sections with a smooth transition. The outer cover may be a material of suitable flexibility and compatibility such as a polyurethane, low density polyethylene, Pebax, or Nylon. The exterior and interior surfaces of the catheter or catheter section may be coated with a lubricious layer such as a hydrophilic polymer layer, which is either chemically bonded to the layer or is physically coated on the surface.

The inner liner 126 may be omitted, particularly in the more proximal segment 124 since the majority of materials which are suitable for the more proximal segment 124 are very hard and suitably slippery for passage of guidewires and the like. The more proximal segment 124 may be a simple tubular member comprising unfilled, filled, or fiber-reinforced, tough, polymeric materials preferably having high flexural moduli. Examples include polyamides, polyamide-polyimides, polyimides (thermoset and thermoplastic), polycarbonates, LCP's, and acetals, for example. To integrate the more proximal section of the catheter assembly with materials found in adjacent sections, the choice of materials for the proximal section is desirably a polyamide which is melt-miscible with a polymeric component found in the adjacent more distal section. The distal section may have a covering of polyurethane, a block copolymer of a polyether and a polyamide (e.g., Pebax), or a low durometer Nylon.

The proximal segment 124 may also comprise a stiffener interposed between an inner liner and an outer cover. The inner liner and outer cover may comprise the materials specified herein for the inner liner 126 and outer cover 130 of the distal segment 122. The outer cover and inner liner of the proximal section may be formed from the same material as the outer cover and inner liner of the distal section, or from different materials. The stiffener may be a knit, braided or coil member comprising a metal alloy such as nitinol or stainless steel, or a polymeric material. The knit stiffener from the distal section may extend into the proximal section. The braid may be formed from a single wire or multiple wires and may comprise more than one material. The coil may be formed from a helically wound wire, having a circular or rectangular cross-section (i.e., ribbon), for example.

In view of the above, it will be seen that the several objects of the invention are achieved and other advantageous results attained.

As various changes could be made in the above constructions and methods without departing from the scope of the invention, it is intended that all matter contained in the above description and shown in the accompanying drawings shall be interpreted as illustrative and not in a limiting sense.

CLAIMSWHAT IS CLAIMED IS:

5 1. A catheter section comprising an elongate tubular member having a proximal end, a distal end, and a passageway defining a lumen extending between the proximal and distal ends, said elongate tubular member comprising a knit tubular member and an inner tubular liner in coaxial relationship with the knit tubular member.

10 2. The catheter section of claim 1 wherein the knit tubular member comprises a metal alloy.

15 3. The catheter section of claim 1 wherein the knit tubular member comprises a superelastic alloy.

20 4. The catheter section of claim 3 wherein the superelastic alloy is a nickel-titanium alloy.

25 5. The catheter section of claim 3 wherein the superelastic alloy is nitinol.

30 6. The catheter section of claim 1 wherein the knit tubular member comprises stainless steel.

 7. The catheter section of claim 1 wherein the knit tubular member comprises a platinum alloy.

 8. The catheter section of claim 1 wherein the knit tubular member comprises a non-metallic material.

9. The catheter section of claim 8 wherein the non-metallic material is a polymeric material.

5 10. The catheter section of claim 1 wherein the knit tubular member comprises a multifilament wire

11. The catheter section of claim 10 wherein the multifilament wire comprises stainless steel and platinum.

10 12. The catheter section of claim 10 wherein the multifilament wire comprises material selected from the group consisting of stainless steel, platinum, and nitinol.

15 13. The catheter section of claim 1 wherein the knit tubular member is formed from wire having a generally circular cross-sectional shape.

14. The catheter section of claim 13 wherein the wire has a diameter of about 0.3 mil - 1.5 mil.

20 15. The catheter section of claim 1 wherein the knit tubular member comprises a first strand made from a first material and a second strand made from a second material.

25 16. The catheter section of claim 1 further comprising an outer tubular cover extending over the knit tubular member.

30 17. The catheter section of claim 16 wherein the outer tubular cover comprises a material selected from the group consisting of polyimide, polyamide, polyethylene, polypropylene, polyvinylchloride, fluoropolymers including PTFE, FEP, Nylon, polyether block amide, vinylidene fluoride, and their mixtures, alloys, copolymers, and block copolymers.

18. The catheter section of claim 16 wherein the outer tubular cover comprises a polymer which can be heat-shrunk onto the knit tubular member.

5 19. The catheter of claim 16 wherein the outer tubular cover is extruded onto the knit tubular member.

20. The catheter of claim 16 wherein the outer tubular cover is bonded onto the knit tubular member.

10 21. The catheter section of claim 16 wherein at least one of the inner tubular liner and the outer tubular cover are radiopaque.

15 22. The catheter section of claim 1 wherein the inner tubular liner comprises a material selected from the group consisting of polyethylene, fluoreopolymer, Nylon, polyether block amide, polyvinyl chloride (PVC), ethyl vinyl acetate (EVA), polyethylene terephthalate (PET), and their mixtures, alloys, and copolymers.

20 23. The catheter section of claim 1 wherein the knit tubular member is generally not radially expandable.

25 24. A catheter section comprising an elongate tubular member having a proximal end, a distal end, and a passageway defining a lumen extending between the proximal and distal ends, said elongate tubular member comprising an inner liner, an outer cover, and a knit tubular member formed from a metal wire.

30 25. The catheter section of claim 24 wherein the metal wire comprises a superelastic alloy.

26. The catheter section of claim 25 wherein the superelastic alloy is nitinol.

27. The catheter section of claim 24 wherein the metal wire has a generally circular cross-section.

28. The catheter section of claim 24 wherein the metal wire is a multifilament wire.

29. The catheter section of claim 28 wherein the multifilament wire comprises stainless steel and platinum.

30. The catheter section of claim 28 wherein the multifilament wire comprises material selected from the group consisting of stainless steel, platinum, and nitinol.

31. A catheter comprising an elongate tubular member having a proximal end, a distal end, and a passageway defining a lumen extending between those ends, said elongate tubular member comprising:

a relatively stiff proximal segment; and

a relatively flexible distal segment comprising a knit tubular member and an inner tubular liner in coaxial relationship with the knit tubular member.

32. The catheter of claim 31 wherein the knit tubular member comprises a superelastic alloy.

33. The catheter of claim 32 wherein the superelastic alloy is nitinol.

34. The catheter of claim 31 wherein the knit tubular member comprises stainless steel.

5 35. The catheter of claim 31 wherein the knit tubular member comprises a platinum alloy.

36. The catheter of claim 31 wherein the knit tubular member comprises a non-metallic material.

10 37. The catheter of claim 36 wherein the non-metallic material is a polymeric material.

38. The catheter of claim 31 wherein the knit tubular member comprises a multifilament wire.

15 39. The catheter of claim 38 wherein the multifilament wire comprises material selected from the group consisting of stainless steel, platinum, and nitinol.

20 40. The catheter of claim 31 wherein the knit tubular member is formed from wire having a generally circular cross-sectional shape.

41. The catheter of claim 40 wherein the wire has a diameter of about 0.3 mil. - 1.5 mil.

25 42. The catheter of claim 31 wherein the knit tubular member comprises a first strand made from a first material and a second strand made from a second material.

30 43. The catheter of claim 31 further comprising an outer tubular cover extending over the knit tubular member.

44. The catheter of claim 43 wherein the outer tubular cover comprises a material selected from the group consisting of polyimide, polyamide, polyethylene, polypropylene, polyvinylchloride, Nylon, polyether block amide, fluoropolymers including PTFE, FEP, low density polyethylene, vinylidene fluoride, and their mixtures, alloys, copolymers, and block copolymers.

45. The catheter of claim 43 wherein the outer tubular cover comprises a polymer which can be heat-shrunk onto the knit tubular member.

46. The catheter of claim 43 wherein the outer tubular cover is extruded onto the knit member.

47. The catheter of claim 43 wherein the outer tubular cover is bonded on the knit member.

48. The catheter of claim 43 wherein at least one of the inner tubular liner and the outer tubular cover are radiopaque.

49. The catheter of claim 31 wherein the knit tubular member is generally not radially expandable.

50. The catheter of claim 31 wherein the proximal segment has an inner proximal liner and an outer proximal cover.

51. The catheter of claim 50 wherein the proximal segment further comprises a braid interposed between the inner proximal liner and the outer proximal cover.

52. The catheter of claim 50 wherein the proximal segment further comprises a coil interposed between the inner proximal liner and the outer proximal cover.

5 53. The catheter of claim 31 wherein the knit tubular member extends into the proximal segment.

CATHETER WITH KNIT SECTION

ABSTRACT OF THE DISCLOSURE

5 A catheter section comprising an elongate tubular member having a proximal end, a distal end, and a passageway defining a lumen extending between the proximal and distal ends. The elongate tubular member includes a knit tubular member and an inner tubular liner in coaxial relationship with the knit tubular member. The catheter section is for use as a distal section in a catheter having a relatively stiff proximal section.

1/2

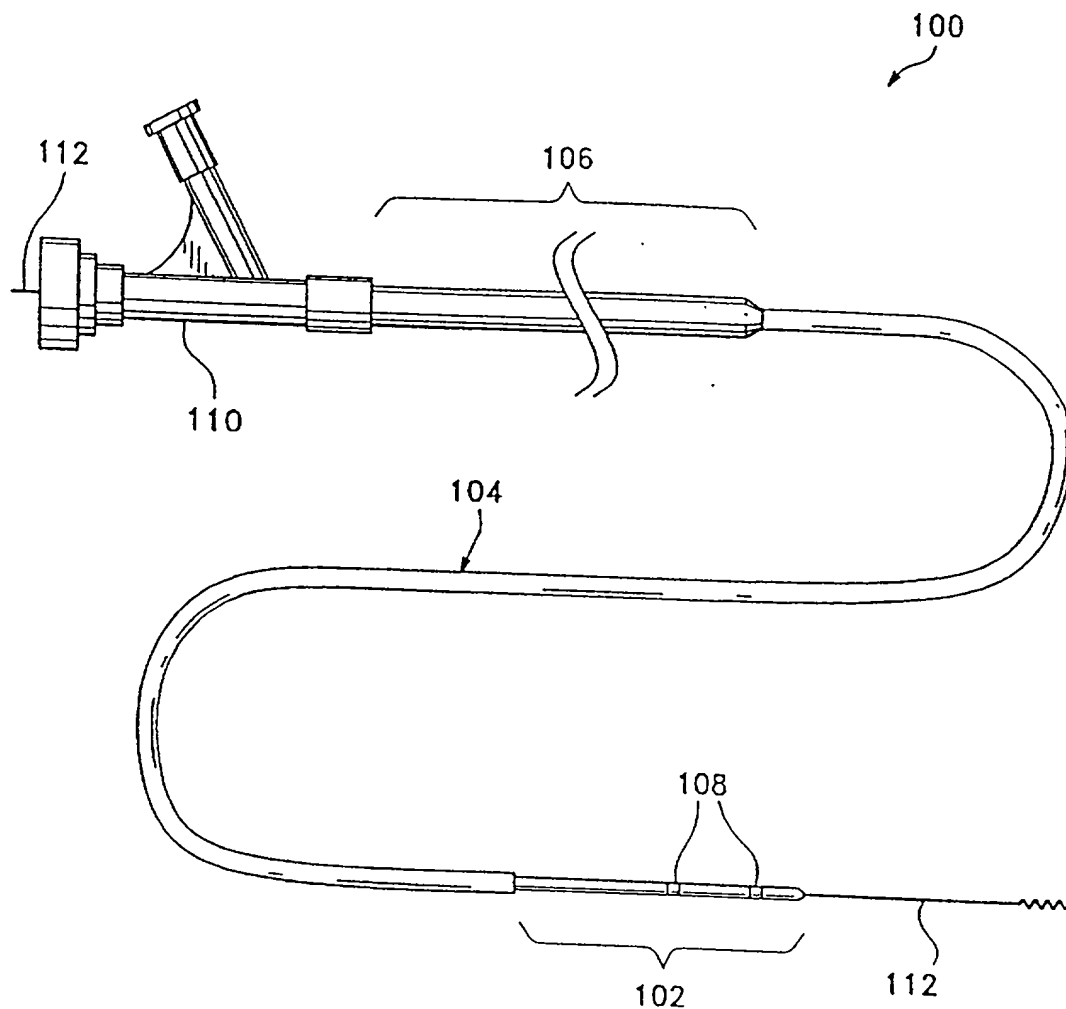
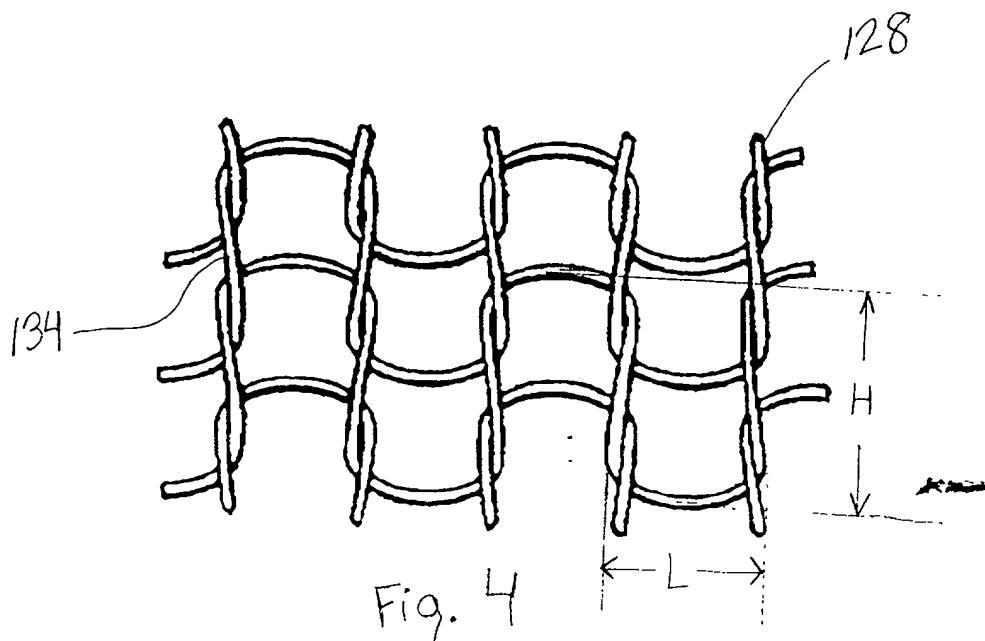


Fig. 1



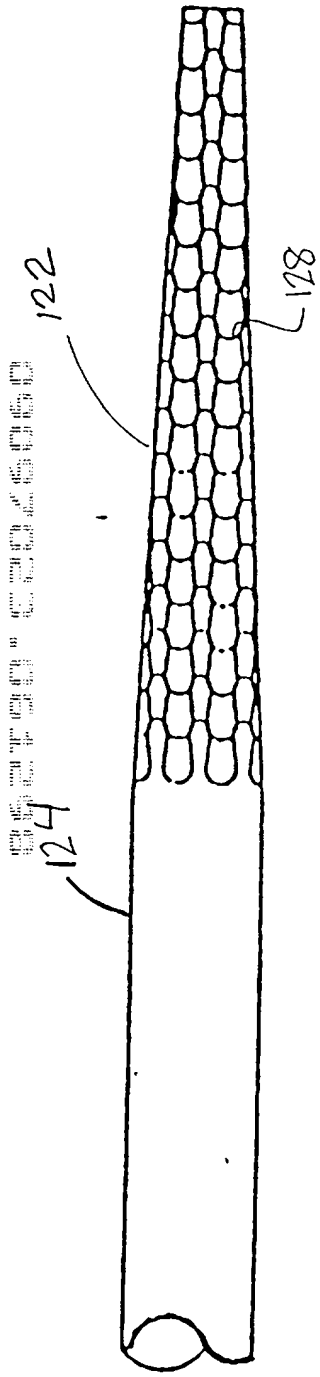


Fig. 2

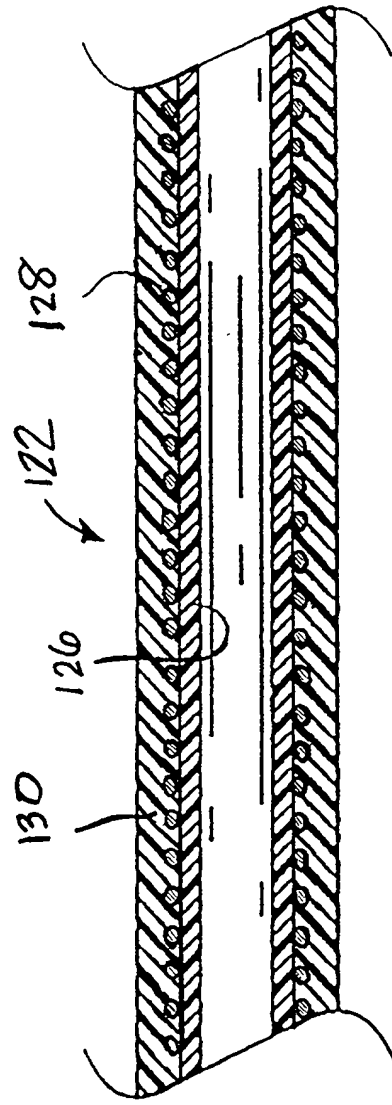


Fig. 3

DECLARATION FOR UTILITY PATENT APPLICATION

AS A BELOW-NAMED INVENTOR, I HEREBY DECLARE THAT:

My residence, post office address, and citizenship are as stated below next to my name.

I believe I am the original, first and joint inventor of the subject matter which is claimed and for which a patent is sought on the invention entitled: CATHETER WITH KNIT SECTION, the specification of which is attached hereto unless the following box is checked:

☐ was filed on _____ as United States Application Serial No. or PCT International Application No. To Be Assigned and was amended on _____ (if applicable).

I HEREBY STATE THAT I HAVE REVIEWED AND UNDERSTAND THE CONTENTS OF THE ABOVE-IDENTIFIED SPECIFICATION, INCLUDING THE CLAIMS, AS AMENDED BY ANY AMENDMENT REFERRED TO ABOVE.

I acknowledge the duty to disclose information which is material to the patentability as defined in 37 C.F.R. § 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT International application which designated at least one country other than the United States listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed:

Application No.	Country	Date of Filing (day/month/year)	Priority Claimed?
			<input type="checkbox"/> Yes <input type="checkbox"/> No

I hereby claim benefit under 35 U.S.C. § 119(e) of any United States provisional application(s) listed below:

Application Serial No.	Filing Date

I hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s), or § 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 C.F.R. § 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

Application Serial No.	Filing Date	Status
		<input type="checkbox"/> Patented <input type="checkbox"/> Pending <input type="checkbox"/> Abandoned

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under § 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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